- 13. (New) A method for the preparation of heterogenised catalyst component comprising:
 - a) providing a halogenated precursor component characterized by the formula:

$$X [CH2]n CH3$$
 (I)

wherein X is an halogen and n is an integer within the range of 1-12;

b) reacting the halogenated precursor with an ionic liquid precursor IL to prepare an ionic liquid of the formula:

$$IL^{+}X$$

c) mixing in a solvent the ionic liquid IL⁺X with a metallocene catalyst component of the formula:

$$R''(Cp) (Cp') M Q_2$$
 (II)

wherein:

Cp and Cp' are each independently a substituted or unsubstituted cyclopentadienyl group, M is a metal from Group 4 of the Periodic Table, R" is a structural bridge imparting stereorigidity between Cp and Cp' and Q is a halogen or an alkyl having from 1 to 12 carbon atoms

wherein the amounts of ionic liquid and catalyst component are in a molar ratio (ionic liquid)/(catalyst component) of from 5:1 to 1:5;

d) heterogeniseng the ionic liquid/metallocine system of subparagraph c) by addition of an apolar solvent to induce a precipitation reaction; and

- e) retrieving a metallocene catalyst component heterogenised by said ionic liquid.
- 14. (New) The method of claim 13 wherein the ionic liquid precursor is an N hydrocarbyl imidazole or pyridine.
- 15. (New) The method of claim 14 wherein the ionic liquid and the catalyst component are mixed in approximately equal stoichiometric amounts.
- 16. (New) The method of claim 14 wherein said ionic liquid precursor is an N-R imidazole in which R is an aryl group or an alkyl group having from 1-12 carbon atoms.
- 17. (New) The method of claim 14 wherein the ionic liquid precursor is 1-methy-3-pentylimidazolium bromide or N-pentyl pyridinium bromide.
- 18. (New) The method of claim 13 further comprising prior to subparagraph c) reacting said ionic liquid with an ionic compound characterized by the formula C⁺A⁻ wherein C⁺ is a cation selected from the group consisting of K⁺, Na⁺, NH₄⁺, and A⁻ is an anion selected from the group consisting of PF₆⁻, SbF₆⁻, BF₄⁻, (CF₃-SO₂)N⁻, C1O₄⁻, CF₃-SO₃)₂N⁻, C1O₄⁻, CF₃ SO₃⁻, NO₃⁻ and CF₃CO₂⁻.
- 19. (New) The method of claim 13 wherein the solvent of subparagraph c) is selected from a group consisting of tetrahydrofuron, methylene dichloride, and toluene.

- 20. (New) The method of claim 19 wherein said apolar solvent is a liquid alkane.
- 21. (New) The method of claim 20 wherein said apolar solvent is n-heptane.
- 22. (New) The method of claim 19 further comprising subsequent to subparagraph c) and prior to subparagraph d) evaporating at least a portion of said solvent prior to the addition of said apolar solvent.
- 23. (New) The method of claim 13 wherein the ligand structure of said metallocene catalyst component incorporates a substituted or unsubstituted bis-indenyl ligand structure, a substituted or unsubstituted bis-benzindenyl ligand structure, or a substituted or unsubstituted bis-tetrahydroindenyl ligand structure.
- 24. (New) The method of claim 23 wherein said metallocene catalyst component is an ethylene bis-tetrahydroindenyl zirconium dichloride, dimethyl silyl bis (2-methylbenzindenyl zirconium dichloride, or dimethyl silyl (2-methyl-4-phenyl-indenyl zirconium dichloride.)
- 25. (New) A heteroginized metallocene catalyst component produced by the method of claim 13.

- 26. (New) A heteroginized catalyst system comprising the catalyst component of claim 25 and an activating agent.
- 27. (New) The catalyst system of claim 26 wherein the activating agent is methylaluminoxane and Q is halogen.
- 28. (New) The catalyst system of claim 27 wherein the methylaluminoxane is present in an amount to provide an Al/M ratio within the range of 100 to 1,000.

- 29. (New) A method for the preparation of an alpha olefin polymer comprising:
 - a) providing a heteroginized catalyst system comprising a heteroginized catalyst component produced by the process of claim 13 and an activating agent for said catalyst component;
 - b) introducing said heterogenised catalyst system in an apolar solvent and an alpha olefin monomer into a polymerization reactor;
 - c) operating said reactor under polymerization conditions; and
 - d) recovering an alpha olefin polymer product from said reactor.
- 30. (New) The method of claim 29 wherein said alpha olefin monomer comprises ethylene or propylene.
 - 31. (New) The method of claim 30 wherein said apolar solvent is n-heptane.
- 32. (New) The method of claim 25 wherein said activating agent is methylalumoxane and wherein said ionic liquid precursor is 1-methyl-3-pentylimidazolium bromide or N-pentyl pyridinium bromide.